

Product Data Sheet

Product Name: Ophthalmic acid TFA

Cat. No.: GC81579

Chemical Properties

Cas. No.

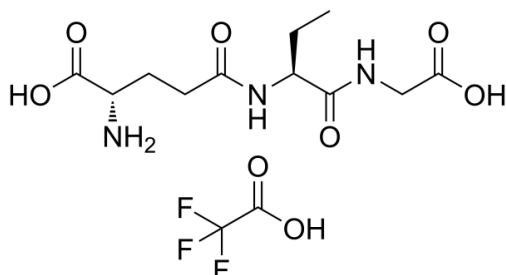
Formula $C_{13}H_{20}F_3N_3O_8$ M.Wt 403.31

Solubility DMSO: 100 mg/mL (247.95 mM; Need ultrasonic) Storage Store at -20°C, away from moisture

General tips For obtaining a higher solubility , please warm the tube at 37 °C and shake it in the ultrasonic bath for a while. Stock solution can be stored below -20°C for several months.

Shipping Condition Evaluation sample solution : ship with blue ice All other available size: ship with RT , or blue ice upon request.

Structure



Caution: Product has not been fully validated for medical applications. For research use only.

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Background

Ophthalmic acid TFA is a ubiquitous metabolite and glutathione modulator, with a K_i of 0.95 mM for glyoxalase I. Ophthalmic acid TFA competitively inhibits glyoxalase I, glutathione S-transferase, glutaredoxin, glutamate-cysteine ligase, protein disulfide reductase (glutathione), as well as non-selective cation channels. Ophthalmic acid TFA is applicable in diabetes-related research [1] [2].

In Vivo, Ophthalmic acid TFA increases liver Ophthalmic acid TFA levels >35-fold, elevates serum Ophthalmic acid TFA, and enhances adipose tissue glucose uptake, while leaving Ophthalmic acid TFA levels unchanged in other tissues in wild-type mice under dietary methionine restriction^[1]. Ophthalmic acid TFA elevates liver Ophthalmic acid TFA levels 20-fold while depleting hepatic glutathione in Cystathionine β -synthase-deficient mice with genetic homocystinuria^[1].

In Vitro, Ophthalmic acid (5 μ M-millimolar concentrations) TFA exerts pleiotropic effects on GSH utilizing enzymes: it induces glutathione reductases, inhibits glutaredoxins (60% reduction at 5 μ M) and millimolar-concentration exposed glutathione S-transferases, and does not affect glutathione peroxidases^[1]. Ophthalmic acid TFA competitively inhibits the glyoxalase I reaction^[1]. Ophthalmic acid TFA inhibits the activation of non-selective cation channels by intracellular NAD^+ via mimicry of GSH's tripeptide structure^[1]. Ophthalmic acid TFA competitively inhibits GSH uptake in yeast^[1]. Ophthalmic acid TFA competitively inhibits purified glyoxalase I enzyme activity with an inhibitor equilibrium constant (K_i) of 0.95 mM^[2].

References:

[1]. Schomakers BV, et al. Ophthalmic acid is a glutathione regulating tripeptide. FEBS J. 2024;291(15):3317-3330.

[2]. CLIFFE EE, et al. The mechanism of the glyoxalase I reaction, and the effect of ophthalmic acid as an inhibitor. Biochem J. 1961 Jun;79(3):475-82.

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